What is claimed is:

```
A DNA sequence encoding a polypeptide of the
1
2
    formula
3
                               WYBAZCX
            wherein WYBAZCX is composed of the polypeptide
    segments shown in Figure 31 (SEQ ID Nos. 136-139, 141-147,
    160, 161, and 163); wherein W comprises polypeptide segment
    F, or is absent; wherein Y comprises polypeptide segment E,
7
    or is absent; wherein Z comprises polypeptide segment G or
    is absent; and wherein X comprises polypeptide segments C/D
 9
    HKL, C/D H, C/D HL, C/D D, C/D' HL, C/D' HKL, C/D' H, C/D'
10
    D, C/D C/D' HKL, C/D C/D' H, C/D C/D'/ HL, C/D C/D' D, C/D D'
11
    H, C/D D' HL, C/D D' HKL, C/D' D' H, C/D' D' HKL, C/D C/D'
12
    D' H, C/D C/D' D' HL, C/D C/D'\D' HKL, or C/D' D' HL;
13
    provided that, either
14
15
            a) at least one of F, Y, B, A, Z, C, or X is of
16
    bovine origin; or
            b) Y comprises polypeptide segment E; or
17
18
            c) X comprises polypeptide segments C/D HKL, C/D D,
    C/D' HKL, C/D C/D' HKL, C/D C/D' D, C/D D' H, C/D D' HL, C/D
19
    D' HKL, C/D' D' H, C/D' D' HKL, C/D C/D' D' H, C/D C/D' D'
20
    HL, C/D C/D' D' HKL, C/D'H, C/D C/D'H, or C/D C/D'HL.
21
 1
            2.
                 The DNA sequence of claim 1, wherein X
    comprises polypeptide segments C/D HKL having the amino acid
 2
    sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-142,
     146, 147, 160, 161).
```

The DNA sequence of claim 1, wherein X 1 comprises polypeptide segments C/D' H having the amino acid 2 sequences shown in Pigure 31 (SEQ ID Nos. 136-139, 141, 143, 3 146, 160). 4 The DNA sequence of claim 1, wherein X 1 comprises polypeptide segments C/D D having the amino acid 2 sequences shown in Figure 31 (SEQ ID Nos. 136/139, 141, 142, 3 144, 160). The DNA sequence of claim 1, wherein X 1 comprises polypeptide segments C/D' HKL/having the amino 2 acid sequences shown in Figure, 31 (SEQ/ID Nos. 136-139, 141, 143, 146, 147, 160, 161). The DNA sequence of claim 1, wherein X 6. 1 comprises polypeptide segments C/D C/D' HKL having the amino 2 acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-3 143, 146, 147, 160, 161). The DNA sequence of claim 1, wherein X 1 comprises polypeptide segments C/D C/D' H having the amino 2 acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-3 143, 146, 160). The DNA sequence of claim 1, wherein X 1 comprises polypeptide segments C/D C/D' HL having the amino

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1 1

acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-

143, 146, 147, 160).

- 9. The DNA sequence of claim 1, wherein X comprises polypeptide segments C/D C/D' D having the amino acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-44, 160).
- 1 10. The DNA sequence of claim 1, wherein X
 2 comprises polypeptide segments C/D D'H having the amino acid
 3 sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-142,
 4 145, 146, 160).
- 11. The DNA sequence of claim 1, wherein X
 2 comprises polypeptide segments C/D D'H L having the amino
 3 acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 1414 142, 145, 146, 147, 160).
- 1 12. The DNA sequence of claim 1, wherein X comprises polypeptide segments C/D D'H K L having the amino 3 acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-42, 145-147, 160, 161).
- 1 13. The DNA sequence of claim 1, wherein X comprises polypeptide segments C/D' D' H having the amino acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141, 143, 145, 146, 160).
- 14. The DNA sequence of claim 1, wherein X
 2 comprises polypeptide segments C/D' D' H K L having the
 3 amino acid sequences shown in Figure 31 (SEQ ID Nos. 1364 139, 141, 143, 145-147, 160, 161).

- 15. The DNA sequence of claim 1, wherein X

 2 comprises polypeptide segments C/D C/D' D' H having the

 3 amino acid sequences shown in Figure 31 (SEQ ID Nos. 136
 4 139, 141-143, 145, 146, 160).
- 1 16. The DNA sequence of claim 1, wherein X 2 comprises polypeptide segments C/D C/D' D' H I having the 3 amino acid sequences shown in Figure 31 (SEQ/ID Nos. 136-139, 141-143, 145-147, 160).
- 1 17. The DNA sequence of claim 1, wherein X 2 comprises polypeptide segments C/D C/D' D' H K L having the 3 amino acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-143, 145-147, 160, 151).
- 18. The DNA sequence comprising coding segments
 2 5'FBA'3' coding for polypeptide segments having the amino
 3 acid sequences shown in Figure 31 (SEQ ID Nos. 136, 138,
 4 139).
- 19. The DNA sequence comprising coding segments
 2 5'FBA'3' coding for polypeptide segments having the amino
 3 acid sequences shown in Figure 31 (SEQ ID Nos. 136, 138,
 4 140).
- 20. The DNA sequence comprising coding segments
 2 5'FEBA' coding for polypeptide segments having the amino
 3 acid sequences shown in Figure 31 (SEQ ID Nos. 136-139,
 4 163).

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21. The DNA sequence comprising coding segments 1 5'FEBA'3' coding for polypeptide segments having the amino 2 acid sequences shown in Figure 31 (SEQ ID Nos. 136-138, 140, 3 163). 22. Purified DNA encoding GGF2HBS5. . 1 23. A polypeptide of the formula/ 1 WYBAZCX 2 wherein WYBAZCX is composed of the polypeptide 3 segments shown in Figure 31 (SEQ ID Nos. 13/6-139, 141-147, 160, 161, 163); wherein W comprises polypeptide segment F, 5 or is absent; wherein Y comprises polypeptide segment E, or 6 is absent; wherein Z comprises polypeptide segment G or is absent; and wherein X comprises peptide segments C/D HKL, 8 C/D H, C/D HL, C/D D, C/D' HL, C/D'/HKL, C/D' H, C/D' D, C/D C/D' HKL, C/D C/D' H, C/D C/D' HL, C/D C/D' D, C/D D' H, C/D 10 D' HL, C/D D' HKL, C/D' D' H, C/p' D' HKL, C/D C/D' D' H, 11 C/D C/D' D' HL, C/D C/D' D' HKJ/, or C/D' D' HL; provided 12 that, either 13 a) at least one of F/, Y, B, A, Z, C, or X is of 14 bovine origin; or 15 b) Y comprises polypeptide segment E; or 16 C) X comprises polypeptide segments C/D HKL, C/D' 17 HKL, C/D D, C/D C/D' HKI, C/D C/D' D, C/D D' H, C/D D' HL, 18 C/D D' HKL, C/D' D' H, C/D' D' HKL, C/D C/D' D' H, C/D C/D' 19 D' HL, C/D C/D' D' HKL, C/D'H, C/D C/D'H, or C/D C/D'HL.

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24. A polypeptide of claim 23, wherein X comprises 1 C/D HKL polypeptide segments having the amino acid sequences 2 shown in Figure 31 (SEQ ID Nos. 136-139, 141-142, 146, 1/47, 3 160, 161). 25. A polypeptide of claim 23, wherein X comprises 1 C/D D polypeptide segments having the amino acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141, 142, 144, 3 160). 26. A polypeptide of claim 23, wherein X comprises 1 C/D' H polypeptide segments having the amino acid sequences 2 shown in Figure 31 (SEQ ID Nos. 136-139, 141, 143, 146, 160). 27. A polypeptide of claim 23, wherein X comprises 1 C/D' HKL polypeptide segments having the amino acid 2 sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141, 143, 146, 147, 160, 161). 4 A polypeptide of claim 23, wherein X comprises 1 C/D C/D' HKL polypeptide segments having the amino acid 2 sequences shown in Figure/31 (SEQ ID Nos. 136-139, 141-143, 3 146, 147, 160, 161). 4 A polypeptide of claim 23, wherein X comprises 1 C/D C/D' H polypeptide segments having the amino acid 2 sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-143, 3 146, 160).

```
30. A polypeptide of claim 23, wherein X comprises
1
   C/D C/D' HL polypeptide segments having the amino acid
2
   sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-
   143,146, 147, 160).
           31. A polypeptide of claim 23, wherein X comprises
1
   C/D C/D' D, polypeptide segments having the amino acid
2
   sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-144,
3
4
    160).
           32. A polypeptide of claim 23, wherein X comprises
1
    C/D D'H polypeptide segments having the amino acid
2
    sequences shown in Figure 31 (SEQ ID Nos./136-139, 141, 142,
3
    145, 146, 160).
            33. A polypeptide of claim 23, wherein X comprises
1
    C/D D'H L polypeptide segments having the amino acid
2
    sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141, 142,
 3
    145-147, 160).
 4
            34. A polypeptide of claim 23, wherein X comprises
 1
    C/D D'H K L polypeptide segments having the amino acid
 2
    sequences shown in Figure 31 /(SEQ ID Nos. 136-139, 141, 142,
 3
     145-147, 160, 161).
                 A polypeptide of claim 23, wherein X comprises
 1
     C/D' D' H polypeptide segments having the amino acid
     sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141, 143,
     145, 146, 160).
```

36. A polypeptide of claim 23, wherein X comprises 1 2 C/D' D' H K L polypeptide segments having the amino acid: sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141, 143, 3 145-147, 160, 161). 4 37. A polypeptide of claim 23, wherein X comprises 1 2 C/D C/D' D' H polypeptide segments having the amino acid sequences shown in Figure 31 (SEQ ID Nos. 136-139, 141-143, 3 4 145, 146, 160). A polypeptide of claim 23, wherein X comprises 1 C/D C/D' D' H L polypeptide segments having the amino acid 2 sequences shown in Figure 31 (SÉQ ID Nos. 136-139, 141-143, 3 4 145-147, 160). 39. A polypeptide of claim 23, wherein X comprises 1 C/D C/D' D' H K L polypeptide segments having the amino acid 2 sequences shown in Figure 31 (SEQ ID/Nos. 136-139, 141-143, 3 145-147, 160, 161). 4 A polypeptide comprising FBA polypeptide 1 segments having the amino acid sequences shown in Figure 31 2 (SEQ ID Nos. 136, 138, 139). 3 41. A polypeptide comprising FEBA polypeptide 1 segments having the amino acid sequences shown in Figure 31 2 (SEQ ID Nos. 136-139, 163). 3 A polypeptide comprising FBA' polypeptide 1 segments having the amino acid sequences shown in Figure 31 2

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(SEQ ID Nos. 136, 139, 140).

3

- 1 43. A polypeptide comprising FEBA' polypeptide 2 segments having the amino acid sequences shown in Figure 31 3 (SEQ ID Nos. 136-139, 140, 163).
- 1 44. Purified GGF2HBS5 polypeptide.
- 45. A basic polypeptide factor having mitogenic activity stimulating the division of Schwann cells in the presence of fetal calf plasma, said polypeptide having a molecular weight of from about 30 kD to about 36 kD, said polypeptide including within its amino acid sequence any one or more of the following polypeptide sequences:

```
FKGDAHTE
7
       ASLADEYEYMXK
8
       TETSSSGLXLK
9
       ASLADEYEYMRK
10
       AGYFAEXAR
11
       TTEMASEQGA
12
       AKEALAALK
13
       FVLQAKK
14
       ETQPDPGQI/LKKVPMVIGAYT
15
       EYKCLKFKWFKKATVM
16
       EXKFYVP
17
       K L E F L X A K,
18
```

46. A basic polypeptide factor having mitogenic activity stimulating the division of Schwann cells in the presence of fetal calf plasma, said polypeptide having a molecular weight of from about 55 kD to about 63 kD, and said polypeptide including within its amino acid sequence any one or more of the following peptide sequences:

VHQVWAAK 7 YIFFMEPEAXSSG 8 LGAWGPPAFPVXY WFVVIEGK 10 ASPVSVGSVQELVQR 11 VCLLTVAALPPT 12 KVHQVWAAK 13 KASLADSGEYNXK 14 DLLLXV 15 EGKVHPQRRGALDRK 16 PSCGRLKEDSK 17 ELNRKNKPQN I KIQKK 18

4

47. A method for stimulating mitogenesis of a glial cell, said method comprising contacting said glial cell with a polypeptide defined by the formula

WYBAZCX

wherein WYBAZCX is composed of the polypeptide 5 segments shown in Figure 31 (SEQ ID Nos. 136-139, 141-147, 6 160, 161, 163); wherein W comprises polypeptide segment F, 7 or is absent; wherein Y comprises polypeptide segment E, or is absent; wherein Z comprises polypeptide segment G or is 9 absent; and wherein X comprises polypeptide segments C/D 10 HKL, C/D H, C/D HL, C/D D, C/D' HL, C/D' HKL, C/D' H, C/D' 11 D, C/D C/D' HKL, C/D C/D' H, C/D C/D' HL, C/D C/D' D, C/D D' 12 H, C/D D' HL, C/D D' HKL, C/D' D' H, C/D' D' HL, C/D' D' 13 HKL, C/D C/D' D' H, C/D C/D' D' HL, or C/D C/D' D' HKL. 14

48. A method for stimulating mitogenesis of a glial 1 cell, said method comprising contacting said glial cell with 2 a polypeptide comprising FBA polypeptide segments having the amino acid sequences shown in Figure 31 (SEQ ID Nos. 136, 138, 139). 5 49. A method of stimulating mitogenesis of a glial 1 cell, said method comprising contacting said glial cell with 2 a polypeptide comprising FBA' polypeptide segments having 3 the amino acid sequences shown in Figure 31/(SEQ ID Nos. 4 136, 138, 140). 5 50. A method of stimulating mitogenesis of a glial 1 cell, said method comprising contacting said glial cell with 2 a polypeptide comprising FEBA polypeptide segments having 3 the amino acid sequences shown in Figure 31 (SEQ ID Nos. 4 136-139, 163). 5 51. A method of stimulating mitogenesis of a glial 1 cell, said method comprising contacting said glial cell with 2 a polypeptide comprising FEBA' polypeptide segments having 3 the amino acid sequences corresponding to polypeptide segments shown in Figure 31/(SEQ ID Nos. 136-138, 140, 163) 5 6 to glial cells. A method of stimulating mitogenesis of a glial cell, said method comprising contacting said glial cell with 1 2 GGF2HBS5 polypeptide. 3 53. A method of stimulating mitogenesis of a glial cell said method comprising contacting said glial cell with 1 a compound which specifically binds the plaserba2 receptor 2 3 of glial cells. 4

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- 54. A method of stimulating mitogenesis of a glial cell, said method comprising contacting said glial cell with a polypeptide, comprising EGFL1, having the amino acid sequence shown Fig. 38, Seq. ID No. 154.

 55. A method of stimulating mitogenesis of a glial cell, said method comprising contacting said glial cell with a polypeptide, comprising EGFL2, having the amino acid sequence shown in Figure 39, Seq. ID No. 155.
- 56. A method of stimulating mitogenesis of a glial cell, said method comprising contacting said glial cell with a polypeptide, comprising EGFL 3, with the amino acid sequence shown in Fig. 40, Seq. ID No. 156.
- 57. A method of stimulating mitogenesis of a glial cell, said method comprising contacting said glial cell with a polypeptide, comprising EGFL4, with the amino acid sequence shown in Fig. 41, Seg. ID No. 157.
- 58. A method of stimulating mitogenesis of a glial cell, said method comprising contacting said glial cell with a polypeptide, comprising EGFL5, with the amino acid sequence shown in Fig. 42, Seq. ID No. 158, to glial cells.
- 59. A method of stimulating mitogenesis of a glial cell, said method comprising contacting said glial cell with a polypeptide, comprising EGFL6, with the amino acid sequence shown Fig. 43, Seq. ID No. 159.
- 1 60. A method for the prophylaxis or treatment of a 2 pathophysiological condition of the nervous system in a 3 mammal in which said condition involves a cell type which is

sensitive or responsive to a polypeptide as defined in any one of claims 1 and 18-22, said method comprising administering to said mammal an effective amount of said polypeptide. 61. A method as claimed in claim 60, wherein said 1 condition involves peripheral nerve damage. 2 The method as claimed in claim 60, wherein said 1 condition involves glia of the central nervous system. 63. A method of stimulating/mitogenic activity in a 1 glial cell, said method comprising/applying 35 kD 2 polypeptide factor isolated from the rat I-EJ transformed fibroblast cell line to said glial cell. A method of stimulating mitogenic activity in a 1 glial cell, said method comprising applying 75 kD 2 polypeptide factor isolated from the SKBR-3 human breast 3 cell line to said glial cell. 4 65. A method of/stimulating mitogenic activity in a 1 glial cell, said method comprising applying 44 kD 2 polypeptide factor isolated from the rat I-EJ transformed fibroblast cell line to said glial cell. 66. A method of stimulating mitogenic activity in a 1 glial cell, said method comprising applying 45 kD 2 polypeptide factor isolated from the MDA - MB 231 human 3 breast cell line to said glial cell.

67. A method of stimulating mitogenic activity in 1 glial cell, said method comprising applying 7 to 14 kD 2 polypeptide factor isolated from the ATL-2 human T-cell line to said glial cell. 68. A method of stimulating mitogenic activity in a 1 glial cell, said method comprising applying 25 AD 2 polypeptide factor isolated from activated mouse peritoneal 3 macrophages to said glial cell. 4 A method of stimulating mitogenic activity in a 1 glial cell, said method comprising applying a 25 kD 2 polypeptide factor isolated from bovine kidney to said glial 3 cell. 4 70. A method of stimulating mitogenic activity in a 1 glial cell, said method comprising applying ARIA polypeptide 3 to said glial cell. A polypeptide factor having glial cell 1 mitogenic activity and including an amino acid sequence 2 encoded by:-3 (a) a DNA sequence shown in any one of Figures 28a, 4 28b or 28c (SEQ ID Nos. 133-135, respectively). 5 (b) a DNA sequence shown in Figure 22 (SEQ ID No. 6 7 89); (c) the DNA sequence represented by nucleotides - B 281-557 of the sequence shown in Figure 28a. 9 a DNA sequence hybridizable to any one of the 10 DNA sequences according to (a), (b) or (c). 11

72. A basic polypeptide factor having a molecular weight, whether in reducing conditions or not, of from about 30 kD to about 36 kD on SDS-polyacrylamide gel electrophoresis, said polypeptide factor having mitogenic activity stimulating the division of rat Schwann cells in the presence of fetal calf plasma, and when isolated using reversed-phase HPLC retaining at least 50% of said activity after 10 weeks incubation in 0.1% trifluoroacetic acid at 4°C.

73. A basic polypeptide factor having a molecular weight, under non-reducing conditions, of from about 55 kD to about 63 kD on SDS-polyacrylamide gel electrophoresis, said polypeptide factor having mitogenic activity stimulating the division of rat Schwann cells in the presence of fetal calf plasma, and when isolated using reversed-phase HPLC retains at least about 50% of said activity after 4 days incubation in 0.1% trifluoroacetic acid at 4°C.

A method for the preparation of a polypeptide 1 defined in claim 72 or claim 73, said method comprising 2 extracting vertebrate brain material to obtain protein, subjecting said protein to chromatographic purification comprising hydroxylapatite HPLC and thereafter to SDS-5 polyacrylamide gel electrophoresis and collecting that 6 fraction therefrom which has an observed molecular weight of 7 about 30 kD to 36 kD and/or that fraction which has an 8 observed molecular weight of about 55 kD to 63 kD if, in 9 either case, subjected to SDS-polyacrylamide gel 10 electrophoresis; in the case of said smaller molecular 11 weight fractions whether in reducing conditions or not, and 12 in the case of said larger molecular weight fraction under 13 non-reducing conditions, and which fraction(s) exhibit(s) 14 mitogenic activity stimulating the division of rat Schwann 15 cells against a background of fetal calf plasma. 16 A method as claimed/in claim 74, wherein the 1 brain material in said method is pituitary material. 2 76. A method as claimed in claim 75, wherein said 1 pituitary material in said/method is bovine. A method as claimed in claim 74, wherein said 1 protein used in said method is initially extracted from 2 brain material is first/subjected to carboxymethyl cellulose chromatography. 4 78. A method as claimed in claim 74 wherein after 1 said hydroxylapatite HPLC, said method uses cation exchange 2 chromatography, gel filtration, and/or reversed-phase HPLC. 3

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79. A method as claimed in claim 74, wherein at 1 each stage of said method biological activity of material obtained is assessed for mitogenic activity stimulating the 3 division of rat Schwann cells in the presence of fetal calf plasma. 5 A method for assaying a substances for glial 1 cell mitogenic activity, said method comprising/contacting 2 said substance with glial cells in the presence of fetal 3 calf plasma, and the measuring DNA synthesis in said glial cells as a measure of glial cell mitogenic activity. 81. An assay as claimed, in claim 80, wherein said glial cells are Schwann cells. A DNA sequence encoding a polypeptide having 1 glial cell mitogenic activity and /comprising: .. (a) a DNA sequence shown in any one of Figures 28a, 3 28b, or 28c (SEQ ID Nos. 133-135) 4 (b) a DNA sequence shown in Figure 22 (SEQ ID No. 5 89); 6 (c) the DNA sequence represented by nucleotides 7 281-557 of the sequence shown in Figure 28a; or 8 (d) a DNA sequence hybridizable to any one of the 9 DNA sequences according to (a), (b) or (c). 10 A polypeptide which is a glial cell mitogen, 1 said polypeptide being encoded by a DNA sequence as defined in claim 82, said polypeptide obtained by a method comprising for the preparation of a glial cell mitogenic factor, said method cultivating modified host cells under conditions permitting expression of said DNA sequence.

84. A vector comprising a DNA sequence as defined 1 in claim 82. 85. A host cell containing the isolated DNA of 1 2 claim 84. 86. A method for the preparation of a glial cell 1 mitogenic factor, said method comprising cultivating 2 modified host cells as defined in claim 85 under conditions permitting expression of said DNA sequence. A polypeptide which is a glial cell mitogen, 87. 1 said polypeptide being encoded by a DNA sequence as defined 2 in claim 1, said polypeptide obtained by a method comprising 3 for the preparation of a glial cell mitogenic factor, said 4 method cultivating modified host cells under conditions permitting expression of said DNA sequence. A polypeptide which is a glial cell mitogen, 1 said polypeptide being encoded by a DNA sequence as defined 2 in any one of claims 18-22, said polypeptide obtained by a method comprising for the preparation of a glial cell mitogenic factor, said method cyltivating modified host cells under conditions permitting expression of said DNA sequence. 7 89. A method for detecting, in a sample, the 1 presence of a molecule having a receptor binding 2 characteristic of a polypeptide defined in any one of claims 3 23, 40-46, 71-73, or 87, said method comprising the steps of contacting said sample with a polypeptide of any 5 one of claims 22, 39-42, 63-65, 72, 73 or 80, along with a

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7 receptor capable of binding specifically to said
8 polypeptide, and
9 b) detecting competitive inhibition of the binding

V 2

- of said polypeptide to said receptor as an indication of the presence of a receptor binding molecule in said sample.
 - glial tumor in a patient, said method comprising
 administering to said patient an effective amount of a
 substance which inhibits the binding of a factor as defined
 in any one of claims 23, 40-46, 71-73, or 87 to a receptor
 therefor.
 - 91. A pharmaceutical or veterinary formulation 2 comprising a polypeptide as defined in any of claims 23, 40-3 46, 71-73, or 87 formulated for pharmaceutical or veterinary 4 use, respectively, together with an acceptable diluent, 5 carrier or excipient and/or in unit dosage form.
 - 1 92. A method for stimulating mitogenesis of a glial cell, said method comprising contacting said glial cell with a polypeptide as defined in any one of claims 23, 40-46, 71-473, or 87.
 - 93. A polypeptide, as defined in any one of claims 2 23, 40-46, 71-73, or 87 for use as a glial cell mitogen.
 - 94. A method for stimulating mitogenesis of a glial cell in a vertebrate, said method comprising contacting said glial cell with an effective amount of a polypeptide defined in any one of claims 23, 40-46, 71-73, or 87 to glial cells.

95. A method for the prophylaxis or treatment of 1 pathophysiological condition of the nervous system in a 2 mammal in which said condition involves a cell type which is sensitive or responsive to a polypeptide as defined in any one of claims 23, 40-46, 71-73, or 87, said method 5 comprising administering an effective amount of said polypeptide. 7 96. A method for the treatment of a condition which 1 involves peripheral nerve damage in a mammal, said method 2 comprising contacting said peripheral merves with an 3 effective amount of a polypeptide, as defined in any one of claims 23, 40-46, 71-73, or 87. 5 97. A method for the prophylaxis or treatment of a 1 condition in a mammal in said condition involves 2 demyelination or damage or loss of Schwann cells, for 3 example a neuropathy of sensory or motor nerve fibers, said 4 method comprising contacting said Schwann an effective amount of a polypeptide, as defined in any one of claims 23, 6 40-46, 71-73, or 87. 7 A method for the prophylaxis or treatment of a 1 neurodegenerative disorder in a mammal, said method 2 comprising contacting glial cells in a mammal with an 3 effective amount of a polypeptide as defined in any one of claims 23, 40-46, 71-73, or 87. 5 A method for inducing neural regeneration 1 and/or repair in a mammal, said method comprising contacting 2 glial cells in a mammal with an effective amount of a 3 polypeptide as defined in any one of claims 23, 40-46, 71-4 5 73, or 87. - 199 -

100. A method of inducing fibroblast proliferation, 1 said method comprising contacting said fibroblasts with a 2 polypeptide, as defined in any one of claims 23, 40-46, 71-3 73, or 87. 101. A method of wound repair in mammals, said 1 method comprising contacting said wound with a polypeptide, 2 as defined in any one of claims 23, 40-46, 7/1-73, or 87. 3 102. A method of making a medicament comprising 1 admixing a polypeptide as defined in any/one of claims 23, 40-46, 71-73, or 87 with a pharmaceutically acceptable 3 carrier. 103. A method for producing an antibody, said method 1 comprising immunizing a mammal with a polypeptide of any one 2 of claims 23, 40-46, 71-73, or 87/2 3 104. A method for detecting, in a sample, the 1 presence of a molecule having /a receptor binding 2 characteristic of a polypeptide defined in any one of claims 3 23, 40-46, 71-73, or 87, said method comprising the steps of 4 contacting said sample with a polypeptide of any 5 one of claims 23, 40-46, 71-73, or 87, along with a receptor 6 capable of binding specifically to said polypeptide, and 7 b) detecting competitive inhibition of the binding 8 of said polypeptide to said receptor as an indication of the 9 presence of a receptor/binding molecule in said sample. 10 105. A method for detecting a receptor which capable 1 of binding to a polypeptide as defined in any one of claims 2 23, 40-46, 71-73, or 87, said method comprising carrying out 3

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affinity isolation on said sample using a said peptide as
  the affinity ligand.
5
         106. A method for the prophylaxis or treatment of a
1
  glial tumor in a patient, said method comprising
2
  administering to said patient an effective amount of a
3
  substance which inhibits the binding of a factor as defined
4
5 in any one of claims 23, 40-46, 71-73, or 87 to/a receptor
  therefor.
         107. A peptide selected from the following:-
1
         FKGDAHTE
2
         ASLADEYEYMXX
3
         TETSSSGLXLK
         ASLADEYEYMRK
5
         AGYFAEXAR
6
         TTEMASEQGA
7
         AKEALAALK
8
         FVLQAKK
9
         ETQPDPGQILK/KVPMVIGAYT
10
         EYKCLKFKWF/KKATVM
11
         EXKFYVP
12
         KLEFLXAK
13
         VHQVWAAK
14
         YIFFMEPEAXSSG
15
         LGAWGPPAFPVXY
16
         WFVVIEGR
17
          A S P V S V G/S V Q E L V Q R
18
          VCLLTVÄALPPT
19
          KVHQVWÄAK
20
          KASLAD/SGEYHXK
21
          DLLLXV
22
```

108. A DNA sequence as shown in any one of Figures 1 2 28a, 28b and 28c (SEQ ID No. 133-135, respectively). 109. A polypeptide encoded by a DNA sequence as 1 defined in claim 108 (SEQ ID Nos. 133-135). 2 110. An antibody to a polypeptide as defined in 1 2 claim 107. 111. A method of investigating, isolating or 1 preparing a glial cell mitogen or gene sequence encoding 2 said glial cell mitogen, said method comprising contacting tissue preparations or samples with an antibody, said 4 antibody prepared as defined in claim 103/. 5 112. A method for isolating a nucleic acid sequence 1 coding for a molecule having glial cell mitogenic activity, 2 said method comprising contacting a/cell containing sample with a glial cell mitogen specific antibody to determine expression of said mitogen in said sample and isolating said 5 nucleic acid sequence from the cells exhibiting said 7 expression. 113. The purified GGF/2 polypeptide comprising the 1 amino acid sequence shown in Fig. 45 herein (SEQ ID No. 2 3 167). 114. A purified GGF2 DNA encoding the GGF2 1 polypeptide whose sequences is shown in Fig. 45 (SEQ ID No. 3 167). 115. A method/for inducing myelination of a neural 1 cell by a Schwann cell, said method comprising contacting - 202 -

- 3 said Schwann cell with a polypeptide of any one of claims
- 4 23, 40-46, 71-73, or 87.
- 1 116. A method for inducing acetylcholine receptor
- 2 synthesis in a cell, said method comprising contacting of
- 3 said cell with a polypeptide of any one of claims 23, 40-46,
- 4 71-73, or 87.
- 1 117. An antibody to a polypeptide as defined in
- 2 claim 23.
- 1 118. An antibody to a polypeptide as defined in
- 2 claim 40.
- 1 119. An antibody to a polypeptide as defined in
- 2 claim 41.
- 1 120. An antibody to a polypeptide as defined in
- 2 claim 42.
- 1 121. An antibody to a polypeptide as defined in
- 2 claim 43.
- 1 122. An antibody to a polypeptide as defined in
- 2 claim 44.
- 1 123. An antibody to a polypeptide as defined in
- 2 claim 45.
- 1 124. An antibody to a polypeptide as defined in
- 2 claim 46.

- 125. An antibody to a polypeptide as defined in 1
- 2 claim 71.
- 126. An antibody to a polypeptide as defined in
- claim 72. 2
- 127. An antibody to a polypeptide as defined in 1
- claim 73.
- 128. An antibody to a polypeptide as defined in 1
- 2 claim 87.
- 129. A method of purifying a protein with glial cell 1
- mitogenic activity, said method comprising contacting a cell 2
- extract with an antibody of any one of claims 117-128.
- 130. A method of treating a mammal suffering from a 1
- disease of glial cell proliferation, said method comprising 2
- administering to said mammal/an antibody of any one of
- claims 117-128.
- 131. A vector comprising a DNA sequence as defined 1
- in any one of claims 1 or 18-22. 2

 $Add B^{2}$ $Add D^{4}$ Add T= 2